

By: Ellis

S.B. No. 1886

A BILL TO BE ENTITLED

AN ACT

relating to diagnostic testing of pregnant women and certain newborns.

BE IT ENACTED BY THE LEGISLATURE OF THE STATE OF TEXAS:

SECTION 1. The heading to Section 81.090, Health and Safety Code, is amended to read as follows:

Sec. 81.090. DIAGNOSTIC [~~SEROLOGIC~~] TESTING DURING PREGNANCY AND AFTER BIRTH.

SECTION 2. Section 81.090, Health and Safety Code, is amended by amending Subsections (a), (b), (c), (i), (j), (k), and (l) and adding Subsections (a-1), (c-1), and (c-2) to read as follows:

(a) A physician or other person permitted by law to attend a pregnant woman during gestation or at delivery of an infant shall:

(1) take or cause to be taken a sample of the woman's blood or other appropriate specimen at the first examination and visit;

(2) submit the sample to an appropriately certified [a] laboratory [~~approved under this section~~] for diagnostic testing approved by the United States Food and Drug Administration for:

(A) [~~a standard serologic test for~~] syphilis [~~approved by the board~~];

(B) [~~a standard serologic test for~~] HIV infection [~~approved by the board~~]; and

1 (C) [~~a standard serologic test for~~] hepatitis B
2 infection [~~approved by the board~~]; and

3 (3) retain a report of each case for nine months and
4 deliver the report to any successor in the case.

5 (a-1) A physician or other person permitted by law to attend
6 a pregnant woman during gestation or at delivery of an infant shall:

7 (1) take or cause to be taken a sample of the woman's
8 blood or other appropriate specimen at an examination in the third
9 trimester of the pregnancy;

10 (2) submit the sample to an appropriately certified
11 laboratory for a diagnostic test approved by the United States Food
12 and Drug Administration for HIV infection; and

13 (3) retain a report of each case for nine months and
14 deliver the report to any successor in the case.

15 (b) A successor is presumed to have complied with this
16 section if the successor in good faith obtains a record that
17 indicates compliance with Subsections (a) and (a-1), if applicable.

18 (c) A physician or other person in attendance at a delivery
19 shall:

20 (1) take or cause to be taken a sample of blood or
21 other appropriate specimen from the mother on admission for
22 delivery; and

23 (2) submit the sample to an appropriately certified
24 [a] laboratory [~~approved under this section~~] for diagnostic testing
25 approved by the United States Food and Drug Administration for:

26 (A) [~~a standard serologic test for~~] syphilis
27 [~~approved by the board~~]; and

1 (B) [~~a standard serologic test for HIV infection~~
2 ~~approved by the board; and~~

3 [~~(C) a standard serologic test for~~] hepatitis B
4 infection [~~approved by the board~~].

5 (c-1) If the physician or other person in attendance at the
6 delivery does not find in the woman's medical records results from
7 the diagnostic test for HIV infection performed under Subsection
8 (a-1), the physician or person shall:

9 (1) take or cause to be taken a sample of blood or
10 other appropriate specimen from the mother;

11 (2) submit the sample to an appropriately certified
12 laboratory for diagnostic testing approved by the United States
13 Food and Drug Administration for HIV infection; and

14 (3) instruct the laboratory to expedite the processing
15 of the test so that the results are received less than six hours
16 after the time the sample is submitted.

17 (c-2) If the physician or other person in attendance at the
18 delivery does not find in the woman's medical records results from a
19 diagnostic test for HIV infection performed under Subsection (a-1),
20 and the diagnostic test for HIV infection was not performed before
21 delivery under Subsection (c-1), the physician or other person in
22 attendance at delivery shall:

23 (1) take or cause to be taken a sample of blood or
24 other appropriate specimen from the newborn child less than two
25 hours after the time of birth;

26 (2) submit the sample to an appropriately certified
27 laboratory for a diagnostic test approved by the United States Food

1 and Drug Administration for HIV infection; and

2 (3) instruct the laboratory to expedite the processing
3 of the test so that the results are received less than six hours
4 after the time the sample is submitted.

5 (i) Before conducting or causing to be conducted a
6 diagnostic [~~standard serologic~~] test for HIV infection under this
7 section, the physician or other person shall advise the woman that
8 the result of a test taken under this section is confidential as
9 provided by Subchapter F, but that the test is not anonymous. The
10 physician or other person shall explain the difference between a
11 confidential and an anonymous test to the woman and that an
12 anonymous test may be available from another entity. The physician
13 or other person shall make the information available in another
14 language, if needed, and if resources permit. The information
15 shall be provided by the physician or another person, as needed, in
16 a manner and in terms understandable to a person who may be
17 illiterate if resources permit.

18 (j) The result of a [~~standard~~] test for HIV infection under
19 Subsection (a)(2)(B), (a-1), (c-1), or (c-2) [~~(c)(2)(B)~~] is a test
20 result for purposes of Subchapter F.

21 (k) Before the [~~blood~~] sample is taken, the health care
22 provider shall distribute to the patient printed materials about
23 AIDS, HIV, hepatitis B, and syphilis. A health care provider shall
24 verbally notify the patient that an HIV test shall be performed if
25 the patient does not object. If the patient objects, the patient
26 shall be referred to an anonymous testing facility or instructed
27 about anonymous testing methods. The health care provider shall

1 note on the medical records that the distribution of printed
2 materials was made and that verbal notification was given. The
3 materials shall be provided to the health care provider by the
4 department [~~Texas Department of Health~~] and shall be prepared and
5 designed to inform the patients about:

6 (1) the incidence and mode of transmission of AIDS,
7 HIV, hepatitis B, and syphilis;

8 (2) how being infected with HIV, AIDS, hepatitis B, or
9 syphilis could affect the health of their child;

10 (3) the available cure for syphilis;

11 (4) the available treatment to prevent
12 maternal-infant HIV transmission; and

13 (5) methods to prevent the transmission of the HIV
14 virus, hepatitis B, and syphilis.

15 (1) A physician or other person may not conduct a diagnostic
16 [~~standard~~] test for HIV infection under Subsection (a)(2)(B),
17 (a-1), or (c-1) [(c)(2)(B)] if the woman objects. A physician or
18 other person may not conduct a diagnostic test for HIV infection
19 under Subsection (c-2) if a parent, managing conservator, or
20 guardian objects.

21 SECTION 3. Subsections (d), (e), (f), and (h), Section
22 81.090, Health and Safety Code, are repealed.

23 SECTION 4. (a) Subsections (a), (c), (i), and (k), Section
24 81.090, Health and Safety Code, as amended by this Act, apply only
25 to a test performed on or after the effective date of this Act. A
26 test performed before the effective date of this Act is covered by
27 the law in effect immediately before the effective date of this Act,

1 and the former law is continued in effect for that purpose.

2 (b) Subsections (a-1), (c-1), and (c-2), Section 81.090,
3 Health and Safety Code, as added by this Act, and Subsections (b),
4 (j), and (l), Section 81.090, Health and Safety Code, as amended by
5 this Act, apply only to a physician or other person attending a
6 pregnant woman during gestation or at delivery of an infant on or
7 after January 1, 2010.

8 SECTION 5. This Act takes effect September 1, 2009.